and net

10/20/2006 10566558b.trn

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LOGINID: SSSPTA1626GMS

PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

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NEWS 1
                Web Page URLs for STN Seminar Schedule - N. America
NEWS 2
                 "Ask CAS" for self-help around the clock
NEWS 3 AUG 09
                INSPEC enhanced with 1898-1968 archive
NEWS 4 AUG 28 ADISCTI Reloaded and Enhanced
NEWS 5 AUG 30 CA(SM)/CAplus(SM) Austrian patent law changes
NEWS 6 SEP 11 CA/CAplus enhanced with more pre-1907 records
NEWS 7 SEP 21 CA/CAplus fields enhanced with simultaneous left and right
                truncation
                CA(SM)/CAplus(SM) display of CA Lexicon enhanced
NEWS 8
        SEP 25
NEWS 9
        SEP 25
                CAS REGISTRY(SM) no longer includes Concord 3D coordinates
NEWS 10
        SEP 25
                CAS REGISTRY(SM) updated with amino acid codes for pyrrolysine
NEWS 11 SEP 28
                CEABA-VTB classification code fields reloaded with new
                classification scheme
NEWS 12 OCT 19
                The Derwent World Patents Index suite of databases on STN will
                be enhanced and reloaded on October 22, 2006
NEWS 13 OCT 19
                LOGOFF HOLD duration extended to 120 minutes
NEWS 14 OCT 19 E-mail format enhanced
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NEWS EXPRESS JUNE 30 CURRENT WINDOWS VERSION IS V8.01b, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 26 JUNE 2006.

NEWS HOURS STN Operating Hours Plus Help Desk Availability
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NEWS IPC8 For general information regarding STN implementation of IPC 8
NEWS X25 X.25 communication option no longer available

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FILE 'HOME' ENTERED AT 10:35:19 ON 20 OCT 2006

=> Uploading THIS COMMAND NOT AVAILABLE IN THE CURRENT FILE 10/20/2006 10566558b.trn

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=> FILE REGISTRY

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

FULL ESTIMATED COST

ENTRY SESSION 0.21 0.21

FILE 'REGISTRY' ENTERED AT 10:35:32 ON 20 OCT 2006
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STRUCTURE FILE UPDATES: 19 OCT 2006 HIGHEST RN 910855-26-4 DICTIONARY FILE UPDATES: 19 OCT 2006 HIGHEST RN 910855-26-4

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TSCA INFORMATION NOW CURRENT THROUGH June 30, 2006

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http://www.cas.org/ONLINE/UG/regprops.html

=>

Uploading C:\Program Files\Stnexp\Queries\10566558b.str

```
chain nodes :
10  11  12  13  14  17
ring nodes :
1  2  3  4  5  6  7  8  9
chain bonds :
5-11  9-10  11-12  11-13  12-14  12-17
ring bonds :
1-2  1-6  2-3  3-4  4-7  5-6  5-9  6-7  7-8  8-9
exact/norm bonds :
5-6  5-9  5-11  11-13  12-14  12-17
exact bonds :
1-2  1-6  2-3  3-4  4-7  6-7  7-8  8-9  9-10  11-12
isolated ring systems :
containing 1 :
```

G1:X

G2:X,OH

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS 11:CLASS 12:CLASS 13:CLASS 14:CLASS 17:CLASS

Stereo Bonds:

10-9 (Single Wedge).

Stereo Chiral Centers:

9 (Parity=Don't Care)

10566558b.trn

Page 3

10:37

10/20/2006 10566558b.trn

Stereo RSS Sets:

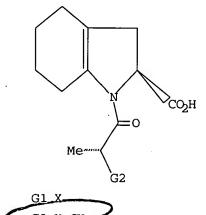
Type=Relative (Default). 1 Nodes= 9

L1 STRUCTURE UPLOADED

=> d 11

L1 HAS NO ANSWERS

L1 STR



Structure attributes must be viewed using STN Express query preparation.

=> s 11

SAMPLE SEARCH INITIATED 10:35:51 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 0 TO ITERATE

100.0% PROCESSED 0 ITERATIONS 0 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 0 TO 0 PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

=> s l1 sss full

FULL SEARCH INITIATED 10:35:57 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 16 TO ITERATE

100.0% PROCESSED 16 ITERATIONS 1 ANSWERS

SEARCH TIME: 00.00.01

L3 1 SEA SSS FUL L1

=> FIL HCAPLUS

COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION

FULL ESTIMATED COST 166.94 167.15

FILE 'HCAPLUS' ENTERED AT 10:36:02 ON 20 OCT 2006

10566558b.trn Page 4 10:37

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FILE COVERS 1907 - 20 Oct 2006 VOL 145 ISS 18 FILE LAST UPDATED: 19 Oct 2006 (20061019/ED)

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s 13.L42 L3

=> d l4 ibib abs hitstr tot

ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2006 ACS on STN 1.4

ACCESSION NUMBER:

2005:1311320 HCAPLUS

DOCUMENT NUMBER:

144:7101

TITLE:

Method for synthesis of perindopril and its

INVENTOR (S):

PATENT ASSIGNEE(S):

SOURCE:

pharmaceutically acceptable salts Fugier, Claude; Dubuffet, Thierry; Langlois, Pascal

Adir et Compagnie, Fr. Eur. Pat. Appl., 9 pp.

CODEN: EPXXDW

DOCUMENT TYPE:

LANGUAGE:

Patent French

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND DATE	APPLICATION NO.	DATE
EP 1367063 EP 1367063	A1 2003/12 B1 200608	203 EP 2003-291931	20030731
R: AT, BE, CH,	DE, DK, ES, F	FR, GB, GR, IT, LI, LU, NL,	
AT 337332	E 200609		20030731
		210 AU 2004-261439	
CA 2533005		CA 2004-2533005	
WO 2005012333	A2 200502	10 WO 2004-FR2035	20040729
WO 2005012333	A3 200503	324	
W: AE, AG, AL,	AM, AT, AU, A	Z, BA, BB, BG, BR, BW, BY,	BZ, CA, CH,
CN, CO, CR,	CU, CZ, DE, D	OK, DM, DZ, EC, EE, EG, ES,	FI, GB, GD,
		L, IN, IS, JP, KE, KG, KP,	
		IA, MD, MG, MK, MN, MW, MX,	
		T, RO, RU, SC, SD, SE, SG,	
		A, UG, US, UZ, VC, VN, YU,	

RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG CN 1826352 Α 20060830 CN 2004-80021209 20040729 BR 2004013169 Α 20061003 BR 2004-13169 20040729 US 2006183920 20060817 A1 US 2006-566562 20060131 NO 2006000922 Α 20060224 NO 2006-922 20060224 PRIORITY APPLN. INFO.: EP 2003-291931 A 20030731 WO 2004-FR2035 W 20040729

OTHER SOURCE(S): MARPAT 144:7101

A method for the synthesis of perindopril [(2S,3aS,7aS)-1-[(2S)-2-[(1S)-1-1]](ethoxycarbonyl)butylamino]propionyl]octahydro-1H-indole-2-carboxylic acid involves coupling of (2S)-hexahydroindole-2-carboxylic acid or its benzyl ester with (R)-G-CHMeCOCl (G = Cl, Br, OH, tosyloxy, mesyloxy or trifluoromethanesulfonyloxy) and then (S)-Et 2-aminopentanoate, followed by catalytic hydrogenation. In an example, the resp. coupling reactions were carried in CH2Cl2-EtNPr-i2 at room temperature and MeCN-Et3N at reflux. Yield of perindopril following hydrogenation was 95% (enantiomeric purity 99%).

IT 870152-15-1P

> RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(synthesis of perindopril from hexahydroindolecarboxylate and bromopropionyl chloride)

RN 870152-15-1 HCAPLUS

CN 1H-Indole-2-carboxylic acid, 1-[(2R)-2-bromo-1-oxopropyl]-2,3,4,5,6,7hexahydro-, (2S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

REFERENCE COUNT:

THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CLATIONS AVAILABLE IN THE RE FORMAT

ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2006 ACS on STN

3

ACCESSION NUMBER:

2005:1311047 **ACAPLUS**

DOCUMENT NUMBER:

144:7100-

TITLE:

Method for synthesis of perindopril and its

INVENTOR(S):

pharmaceutically acceptable salts Fugier, Claude; Dubuffet, Thierry; Langlois, Pascal Adir et Compagnie, Fr.

PATENT ASSIGNEE(S):

SOURCE:

Eur. Pat. Appl., 9 pp.

DOCUMENT TYPE:

CODEN: EPXXDW

Patent

LANGUAGE:

French

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

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PATENT NO.
                                                                       DATE
                                                       KIND
                                                                                                  APPLICATION NO.
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           EP 1367062
                                                                       20031203
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                                                         A 1
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                                                               20060820
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           WO 2005012328
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                            SN, TD, TG
           CN 1826351
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                                                                       20060824
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PRIORITY APPLN. INFO.:
                                                                                                 EP 2003-291930
                                                                                                                                             Α
                                                                                                                                                    20030731
                                                                                                 WO 2004-FR2036
                                                                                                                                             W
                                                                                                                                                    20040729
OTHER SOURCE(S):
                                                       CASREACT 144:7100; MARPAT 144:7100
          A method for the synthesis of perindopril [(2S,3aS,7aS)-1-[(2S)-2-[(1S)-1-(2S)-2-[(1S)-1-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2S)-2-(2
AB
           (ethoxycarbonyl)butylamino]propionyl]octahydro-1H-indole-2-carboxylic
           acid] involves coupling of (2S)-hexahydroindole-2-carboxylic acid or its
          benzyl ester with (R)-G-CHMeCOCl (G = Cl, Br, OH, tosyloxy, mesyloxy or
           trifluoromethanesulfonyloxy) and then (S)-Et 2-aminopentanoate, followed
          by catalytic hydrogenation. In an example, the resp. coupling reactions
           were carried in CH2Cl2-EtNPr-i2 at room temperature and MeCN-Et3N at reflux.
           Yield of perindopril following hydrogenation was 95% (enantiomeric purity
           99%).
           870152-15-1P
           RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
           (Reactant or reagent)
```

ΙT

(synthesis of perindopril from hexahydroindolecarboxylate and bromopropionyl chloride)

RN 870152-15-1 HCAPLUS

CN1H-Indole-2-carboxylic acid, 1-[(2R)-2-bromo-1-oxopropyl]-2,3,4,5,6,7hexahydro-, (2S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

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3

10/20/2006 10566558b.trn

=> log y

COST IN U.S. DOLLARS

SINCE FILE TOTAL
ENTRY SESSION
FULL ESTIMATED COST

15.28

182.43

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)
SINCE FILE TOTAL ENTRY SESSION

CA SUBSCRIBER PRICE -1.50 -1.50

STN INTERNATIONAL LOGOFF AT 10:36:58 ON 20 OCT 2006